

Eplontersen

Regulations: had been registered in the following countries: United States of America (USA).

Registration number (S.A.): Not available.

Insurance Drug Formulary (S.A.): Not available.

General Information:

Registered Company: Ionis Pharmaceuticals and AstraZeneca.

Regulatory Status: R.X.

Mechanism of Action: Plontersen, an antisense oligonucleotide-GalNAc conjugate, is structurally represented as [insert chemical structure here]. This drug causes degradation of mutant and wild-type transthyretin (TTR) mRNA through binding to the TTR mRNA.

Indication

Approved (Labeled) indication: Polyneuropathy due to amyloidosis, Hereditary transthyretin-mediated.

Dosage Forms: Injection.

Dosing/Administration: 45 mg subcutaneously once monthly.

Dose:

Dose in Renal/Hepatic Failure/Geriatric Dose: No dosage adjustment is needed.

Indicated for pediatrics: Safety and effectiveness not established in pediatric patients.

Pharmacokinetic:

Absorption

- T_{max}: 2 hours.

Distribution

- V_d: 12 L

Cost Analysis

Drugs	Drug classes	Approval Indication	Dose	Cost (American Dollar)	Insurance drug formulary(SCHI)
Eplontersen	Antisense Oligonucleotide	Polyneuropathy due to amyloidosis, Hereditary transthyretin-mediated	45 mg subcutaneously once monthly	Around \$500,000.	Not Covered (24.1.2024)

REFERENCES

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Metabolism

- Hepatic

Excretion

Renal excretion: Less than 1% unchanged antisense oligonucleotide (ASO)

Elimination Half-Life

- SubQ: 3 week

Safety:

Common Adverse Reactions (%): Vomiting and vitamin A deficiency.

Severe/rare adverse Reactions (%) : Atrioventricular block.

Drug Interactions: Not reported yet.

Contraindications: Not available.

Precautions: Reduced serum vitamin A levels have been reported.

Monitoring Requirements: Not available.

Sound-Alikes/ Look-Alikes: Not available.

High Alert: Not available.

Boxed warnings or alerts issue: Not available.

Toxicity if antidote required: Not available.

Storage if there is a particular condition. Store in the original carton refrigerated between (2-8°C).

Patient counseling

- Report symptoms of vitamin A deficiency.
- Side effects may include vomiting, blurred vision and cataracts.